Title (en)

THIAZOLE DERIVATIVES AS A2B ANTAGONISTS

Title (de)

THIAZOLDERIVATE ALS A2B-ANTAGONISTEN

Title (fr)

DERIVES DE THIAZOLE EN TANT QU'ANTAGONISTES DE A2B

Publication

EP 1709036 B1 20080305 (EN)

Application

EP 05706936 A 20050120

Priority

- EP 2005000542 W 20050120
- GB 0401336 A 20040121

Abstract (en)

[origin: WO2005070926A1] Compounds of formula (I) in free or salt form, where Ar is phenyl substituted by one or more substituents selected from halogen, cyano and C1- C8-haloalkyl, or naphthyl, R<1> is hydrogen, phenyl optionally substituted by one or more substituents selected from halogen, cyano, hydroxy, C1-C8-alkyl, C1-C8-haloalkyl, C1-C8-alkoxy, C1-C8-alkoxy-Cl-C8 alkyl, carboxy, C1-C8-alkoxycarbonyl and acyloxy, or R<1> is a 5- or 6- membered monovalent heterocyclic group, R<2> is hydrogen, Cl-C8-alkyl, acyl or -CON(R<3>)R<4>, R<3> and R<4> are each independently hydrogen or C1-Cs-alkyl, or together with the nitrogen atom to which they are attached denote a 5- or 6- membered heterocyclic group, and Y is a pyrimidinyl or pyridazinyl group, optionally substituted by at least one C1-Cs-alkyl, C1-C8-alkoxy, C1-C8-alkylthio, C1-C8-alkyl amino, di(Cl-C8-alkyl) amino or acylamino group. The compounds are useful as pharmaceuticals.

IPC 8 full level

C07D 417/04 (2006.01); **A61K 31/501** (2006.01); **A61K 31/506** (2006.01); **A61P 11/00** (2006.01); **C07D 417/14** (2006.01)

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